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What is claimed is:

- An antisense compound 8 to 30 nucleobases in length targeted to the 5'-untranslated region, translational termination region or 3' untranslated region of a nucleic
 acid molecule encoding focal adhesion kinase, wherein said antisense compound inhibits the expression of said focal adhesion kinase.
- 2. The antisense compound of claim 1 which is an 10 antisense oligonucleotide.
 - 3. The antisense compound of claim 2 wherein the antisense oligonucleotide has a sequence comprising SEQ ID NO: 3, 4, 6, 7, 8, 9, 16, 17, 18, 20 or 23.
- 4. The antisense compound of claim 2 wherein the 15 antisense oligonucleotide comprises at least one modified internucleoside linkage.
 - 5. The antisense compound of claim 4 wherein the modified internucleoside linkage is a phosphorothicate linkage.
- 20 6. The antisense compound of claim 2 wherein the antisense oligonucleotide comprises at least one modified sugar moiety.
 - 7. The antisense compound of claim 6 wherein the modified sugar moiety is a 2'-O-methoxyethyl moiety.
- 25 8. The antisense compound of claim 2 wherein the antisense oligonucleotide comprises at least one modified nucleobase.

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9. The antisense compound of claim 8 wherein the modified nucleobase is a 5-methyl cytosine.

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- 10. The antisense compound of claim 2 wherein the antisense oligonucleotide is a chimeric oligonucleotide.
- 5 11. A pharmaceutical composition comprising the antisense compound of claim 1 and a pharmaceutically acceptable carrier or diluent.
 - 12. The pharmaceutical composition of claim 11 further comprising a colloidal dispersion system.
- 10 13. The pharmaceutical composition of claim 11 wherein the antisense compound is an antisense oligonucleotide.
 - 14. The pharmaceutical composition of claim 11 further comprising a chemotherapeutic agent.
- 15. The pharmaceutical composition of claim 14 wherein 15 the chemotherapeutic agent is 5-fluorouracil.
 - 16. A method of inhibiting the growth of a tumor in an animal comprising administering to said animal an effective amount of the pharmaceutical composition of claim 14.
- 17. A method of inhibiting the expression of focal
 20 adhesion kinase in cells or tissues comprising contacting
 said cells or tissue with the antisense compound of claim 1
 so that expression of focal adhesion kinase is inhibited.
- 18. An antisense compound up to 30 nucleobases in length targeted to the coding region, or start site of a nucleic 25 acid molecule encoding focal adhesion kinase, wherein said antisense compound inhibits the expression of said focal

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adhesion kinase and has a sequence comprising at least an 8 nucleobasic portion of SEQ ID NO: 10, 11, 12, 14, 15, 30, 31 or 33.

- 5 19. The antisense compound of claim 18 which is an antisense oligonucleotide.
 - 20. The antisense compound of claim 19 wherein the antisense oligonucleotide comprises at least one modified internucleoside linkage.
- 10 21. The antisense compound of claim 20 wherein the modified internucleoside linkage is a phosphorothicate linkage.
- 22. The antisense compound of claim 19 wherein the antisense oligonucleotide comprises at least one modified 15 sugar moiety.
 - 23. The antisense compound of claim 22 wherein the modified sugar moiety is a 2'-O-methoxyethyl moiety.
- 24. The antisense compound of claim 19 wherein the antisense oligonucleotide comprises at least one modified 20 nucleobase.
 - 25. The antisense compound of claim 24 wherein the modified nucleobase is a 5-methyl cytosine.
 - 26. The antisense compound of claim 19 wherein the antisense oligonucleotide is a chimeric oligonucleotide.
- 25 27. A pharmaceutical composition comprising the antisense compound of claim 18 and a pharmaceutically acceptable carrier or diluent.

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- 28. The pharmaceutical composition of claim 27 further comprising a colloidal dispersion system.
- 29. The pharmaceutical composition of claim 27 wherein the antisense compound is an antisense oligonucleotide.
- 5 30. The pharmaceutical composition of claim 27 further comprising a chemotherapeutic agent.
 - 31. The pharmaceutical composition of claim 30 wherein the chemotherapeutic agent is 5-fluorouracil.
- 32. A method of inhibiting the growth of a tumor in an animal comprising administering to said animal an effective amount of the pharmaceutical composition of claim 30.
 - 33. A method of inhibiting the expression of focal adhesion kinase in cells or tissues comprising contacting said cells or tissue with the antisense compound of claim
- 15 18 so that expression of focal adhesion kinase is inhibited.
 - 34. A method of treating an animal having a disease or condition associated with focal adhesion kinase comprising administering to said animal a therapeutically or
- 20 prophylactically effective amount of an antisense compound 8 to 30 nucleobases in length targeted to a nucleic acid molecule encoding human focal adhesion kinase wherein said antisense compound inhibits the expression of human focal adhesion kinase.
- 25 35. The method of claim 34 wherein the disease or condition is cancer.
 - 36. The method of claim 35 wherein said cancer is of the breast, colon, mouth or skin.

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- 37. The method of claim 34 wherein said disease or condition is an angiogenic disorder.
- 38. The method of claim 37 wherein said angiogenic disorder is retinal neovascularization.
- 5 39. A method of preventing migration of cells associated with expression of focal adhesion kinase comprising administering to said cells a therapeutically or prophylactically effective amount of an antisense compound 8 to 30 nucleobases in length targeted to a nucleic acid 10 molecule encoding human focal adhesion kinase wherein said antisense compound inhibits the expression of human focal adhesion kinase.
- 40. A method of preventing neovascularization associated with expression of focal adhesion kinase in an animal comprising administering to said animal a therapeutically or prophylactically effective amount of an antisense compound 8 to 30 nucleobases in length targeted to a nucleic acid molecule encoding human focal adhesion kinase wherein said antisense compound inhibits the expression of 20 human focal adhesion kinase.
- 41. A method of treating an animal having a disease or condition associated with focal adhesion kinase comprising administering to said animal a therapeutically or prophylactically effective amount of an antisense compound 25 8 to 30 nucleobases in length targeted to a nucleic acid molecule encoding human focal adhesion kinase in combination with a therapeutically or prophylactically effective amount of a chemotherapeutic agent.
- 42. The method of claim 41 wherein the chemotherapeutic 30 agent is 5-fluorouracil.

- 43. The method of claim 41 wherein the disease or condition is cancer.
- 44. The method of claim 43 wherein said cancer is melanoma.